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Question Paper For

DHA Model Question Paper. by Megha

Baby · February 3, 2015. 1) A client with

COPD is admitted to the medical surgical

unit. To help this client maintain a patent

airway and achieve maximal gas exchange

the nurse should. a)instruct the client to

drink 2L of fluid daily. b)Maintain the

client on bed rest.

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and Answer

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exam to get Dubai Health Authority (DHA) license which is the doorway exam for every healthcare professional who needs to be qualified to get the license. Thus we make available of question papers in Nursing, Pharmacy, Medical Lab Technology, Physiotherapy, General Practitioners, Specialist Doctors, dentistry, specialist dentistry and for other medical professionals.

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DHA Exam Questions & Answers 2020
DHA Exam. In this post we will share Past

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Papers of DHA Exam in MCQ Format.
We will share here few MCQs after these MCQs link will be shared where you will be able to Download complete PDF. You are seeing a 45-year-old obese diabetic woman who reports bilateral lower extremity peripheral edema. In addition to diabetes, she has arthritis,

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MOH/ DHA EXAM QUESTIONS WITH ANSWER Question # 1. A client Mr. PK Shetty who has been receiving urokinase has a large bloody bowel movement. ...

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QUESTIONS: MOH/ DHA EXAM ...

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Question Paper Two 1) The nurse is
teaching a client who has iron deficiency
anemia about foods she should include in
her diet. The nurse determines that the
client understands the dietary
modifications if she selects which of the
following from her menu ?

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issues that every Doctor / Nurses /
Pharmacists / Dentist aide is likely to deal
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helps in gauging your preparation level for the DHA exam, what your shortcomings are, and how much more effort you need to put in.

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1. Fertilization of ovum takes place at:
Ans. Ampulla 2. The most frequent causes of maternal death during child birth: Ans. Hemo...

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patient had duodenum metastatic cancer to
duodenum and pancreas and not affected
to other body aprts. Which level you will
be assessed □-increased blood sugar
level.RECENT DHA QUESTIONS
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fluid to pneumonia patient what you will
assess□skin colour, nail [□]

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explore the importance of EFA and eicosanoids on living organisms.

Organization is around five interrelated themes: examination of the biological function of docosahexaenoic acid at the fundamental level of molecular and cellular research; biosynthesis of PUFA in mammals; types of biological markers that can provide information about the adequacy of EFA intake; role that EFA and eicosanoids play in the development of disease states; and in the nutrition of the fetus and newly born infants, especially those born prematurely. Member price, \$100. Annotation copyright by Book News, Inc., Portland, OR

Ion channels are membrane proteins that regulate the permeability of ions across the cell membrane. The sequential opening of

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different types of ion channels produces action potentials in excitable cells. Action potentials are a way for the body to, for example, transmit signals quickly over a long distance. The KV7 family is an important group of voltage-gated potassium channels. Mutations that cause dysfunction in members of the KV7 family are associated with several forms of disease. Compounds that can activate KV7 channels have previously been shown to work as medical treatments. However, the previously available antiepileptic drug retigabine, has been withdrawn due to adverse effects. Thus, there is a need for further development of compounds that target these channels. PUFA and PUFA analogs have previously been demonstrated to activate KV7.1 through an electrostatic mechanism. This thesis investigates new aspects of the interaction between KV7 channels and PUFA-related

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compounds. The data in this thesis are from human KV7 channels expressed in *Xenopus laevis* oocytes. The currents produced by the channels expressed in the oocytes have been studied using two-electrode voltage clamp. Our aim was to study the mechanism for the activation of KV7 channels by PUFA and PUFA analogs. More specifically, we intended to study why the beta subunit KCNE1 abolishes the activating effect of PUFA on KV7.1 and how PUFAs activate KV7.2 and KV7.3. Additionally, we wanted to study aspects that may affect whether these compounds are viable as medical treatments. For instance, whether these compounds can activate channels containing disease-causing mutations and whether we can improve compound selectivity towards certain KV7 channels. In Paper I, we introduce disease-causing mutations found in patients into KV7.1

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and KCNE1. The characterization showed that these channels had altered biophysical properties compared to wild type channels. A PUFA analog was found to activate and, to a large degree, restore wild type-like biophysical properties in the mutated channels regardless of the localization of the mutation in the channel. In Paper II, we demonstrate why PUFA is unable to activate KV7.1 co-expressed with beta subunit KCNE1. KCNE1 induces a conformational change of KV7.1 that moves the S5-Phelix loop closer to the PUFA binding site. This causes negative charges of the loop to attract protons that reduce local pH at the PUFA binding site. The decreased local pH leads to protonation of PUFA and the PUFAs therefore lose their negative charge. Thus, PUFA cannot activate KV7.1 when it is co-expressed with KCNE1. In Paper III, we study a group of PUFA-related substances,

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endocannabinoids, on KV7 channels. One endocannabinoid, Arachidonoyl-L-Serine (ARA-S), was identified as a potent activator of the neuronal M-channel, comprising KV7.2 and KV7.3 heteromers. We study the activating mechanism of ARA-S in KV7.2 and KV7.3, demonstrating how the activating effect is linked to two parts of the channel protein, one in the voltage sensor domain and the other in the pore domain. ARA-S was also found to activate KV7.1 and KV7.5 but not KV7.4, which instead was inhibited. Retigabine, a compound that activates the M-channel but has a different KV7 subtype selectivity compared to ARA-S, was used in combination with ARA-S to maintain a potent effect on the M-channel while limiting the activation of other KV7 channels. In conclusion, the activating effect of PUFA analogs on KV7 channels may be helpful in the development of

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future drug candidates for diseases such as arrhythmia and epilepsy.

Certain long chain polyunsaturated fatty acids (LCPUFAs) are thought to be essential components of the nutrition of infants, including those prematurely born, in the sense that they cannot be synthesised by the immature organism and must therefore be supplied in the diet. Breast milk contains these substances, but many manufactured infant formulae do not. An absence of dietary LCPUFAs has been thought to affect the development of the brain and retina, possibly leading to abnormalities in cognitive and visual function. Considerable multidisciplinary research has been carried out to investigate

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this proposition. Diets free from LCPUFAs have been compared with supplemented formulae, or with breast milk. The conclusions from this research were critically examined by a group of leading paediatricians, nutritionists, experts in visual science and developmental behavioural scientists at a 'Dobbing Workshop' held in the United States in late February, 1997. Each of the Chapters was precirculated to the whole group, commented on before the Workshop, and then exhaustively discussed. The Chapters and Commentaries which are published here have therefore undergone a more extensive peer-review process than is usually the case.

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